10/656,863 10/05/2005

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FILE COVERS 1907 - 5 Oct 2005 VOL 143 ISS 15 FILE LAST UPDATED: 4 Oct 2005 (20051004/ED)

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=>

Uploading C:\Program Files\Stnexp\Queries\10656863\1st look5.str

$$\begin{array}{c} \text{Cy} \\ \text{H} \\ \text{N} \\ \text{G1} \\ \text{H} \\ \end{array}$$

chain nodes : 10 12 13 15 16 17 ring nodes : 1 2 3 4 5 6 7 chain bonds : 1-18 3-17 4-16 7-10 8-13 10-12 10-15 ring bonds : 1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9 exact/norm bonds : 6-9 8-9 8-13 10-12 10-15 exact bonds : 3-17 4-16 5-7 7-8 7-10 1-18 normalized bonds : 1-2 1-6 2-3 3-4 4-5 5-6 isolated ring systems : containing 1 :

G1:0,S

Match level:
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
12:CLASS 13:CLASS 15:Atom 16:CLASS 17:CLASS 18:CLASS

50 ANSWERS

L1 STRUCTURE UPLOADED

=> D L1

L1 HAS NO ANSWERS

L1 S

G1 0, S

Structure attributes must be viewed using STN Express query preparation.

=> S L1

REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:12:33 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 259 TO ITERATE

100.0% PROCESSED 259 ITERATIONS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 4215 TO 6145 PROJECTED ANSWERS: 1130 TO 2228

L2 50 SEA SSS SAM L1

L3 6 L2

=> S L1 FULL

REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

FULL SEARCH INITIATED 14:12:40 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 5144 TO ITERATE

100.0% PROCESSED 5144 ITERATIONS SEARCH TIME: 00.00.01

1464 ANSWERS

L4 1464 SEA SSS FUL L1

L5 25 L4

=> D IBIB HITSTR 1-25

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L10 ANSWER 19 OF 25 CAPLUS COPYRIGHT 2005 ACS on STN

1992:255518 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 116:255518

TITLE: Synthesis and some reactions of 4-(N-

acetyloxindolylidene) -2-aryl-5(4H) -oxazolones

Kandile, Nadia G.; Abdel Latif, Tahia M.; El Sayed,

II

IV

Univ. Coll. Women, Ain Shams Univ., Cairo, Egypt CORPORATE SOURCE:

SOURCE:

AUTHOR(S):

Revue Roumaine de Chimie (1991), 36(1-3), 245-50

CODEN: RRCHAX; ISSN: 0035-3930

DOCUMENT TYPE:

Journal English

LANGUAGE: OTHER SOURCE(S):

CASREACT 116:255518

GT

GI

III

Oxazolones I (R = Ph, Et) were prepared by the condensation of isatin with AB acylglycines RCONHCH2CO2H under Perkin conditions. The hydrolysis of I gave the corresponding deacetyl derivs. The hydrazinolysis of I with hydrazine at room temperature gave cyclic hydrazides II. The aminolysis of I with H2NR1 (R1 = NHPh, CH2Ph, Ph) gave amides III. The cyclization of III in boiling 6N HCl gave lactams IV.

CAPLUS COPYRIGHT 2005 ACS on STN L10 ANSWER 20 OF 25

1991:81699 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 114:81699

Reaction of 2-aryl-4-(N-acetyloxindolylidene)oxazolin-TITLE:

5-ones with B-substituted amines

Kandile, Nadia G. AUTHOR(S):

Univ. Coll. Women, Ain Shams Univ., Heliopolis, Egypt CORPORATE SOURCE:

SOURCE: Oriental Journal of Chemistry (1989), 5(2), 139-42

CODEN: OJCHEG; ISSN: 0970-020X

DOCUMENT TYPE: Journal

English LANGUAGE:

OTHER SOURCE(S): CASREACT 114:81699

$$X = 0$$
 $X = 0$
 $X =$

AB Oxazolinones I (R = Ph, Me, X = X1) reacted with o-(H2N)2C6H4 to give RCONHC(:X1)CONHC6H4NH2-2 which were cyclized by AcONa-AcOH to benzimidazoles II. Treatment of I with o-(H2N)2C6H4 and AcONa gave imidazoles III which were cyclized to the benzimidazoles by AcONa-AcOH. Similar results were obtained with o-H2NC6H4CO2H.

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L10 ANSWER 24 OF 25 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1968:443798 CAPLUS

DOCUMENT NUMBER: 69:43798

TITLE: Indole derivatives

INVENTOR(S): Teshiqawara, Takashi; Kobayashi, Goro; Matsuda,

Yoshiro

SOURCE: Jpn. Tokkyo Koho, 9 pp.

CODEN: JAXXAD

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 42024899 B4 19671129 JP 19651227

GI For diagram(s), see printed CA Issue.

AB A mixture of 2 g. 3-[bis(methylthio)methylene]-2-oxindole, 10 ml. EtOH, and 1.5 g. ethylenediamine is heated 1 hr. to give 50% I [R = H, (R1R2 =) NHCH2CH2NH, pale yellow, m. >296°. Similarly prepared are the following I (R, R1, R2, and m.p. given): Me, (R1R2 =) NHCH2CH2NH, 256°; H, (R1R2 =) NHCH2CH2O, 261-3°; Me, (R1R2 =) NHCH2CH2O, 242°; Me, MeS, EtO2-CCH2NH, 133.5°; Me, MeS, H2NC(:NH)NH, 236°; H, NH2NH, NH2NH, 183-4°; Me, MeS, cyclohexylamino, 90-1°; Me, Bu2N, NH2NH, 168-9°; H, MeS, PhNH, 167-8°; Me, MeS, PhNH, 124-8°; H, PhNH, 212-13°; H, MeS, morpholino, 234-6°; H, morpholino, morpholino, 300°; Me, MeS, MeS, 174-6°; Me, 1-piperidinyl, NH2NH, 188-90°. The products are bactericides and anti-virus drugs.